FILE 'HOME' ENTERED AT 21:58:42 ON 15 MAR 2009

=> fil .bec

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION

0.22

0.22

FULL ESTIMATED COST

FILES 'MEDLINE, SCISEARCH, LIFESCI, BIOTECHDS, BIOSIS, EMBASE, HCAPLUS, NTIS, ESBIOBASE, BIOTECHNO, WPIDS' ENTERED AT 21:59:18 ON 15 MAR 2009 ALL COPYRIGHTS AND RESTRICTIONS APPLY. SEE HELP USAGETERMS FOR DETAILS.

11 FILES IN THE FILE LIST

=> s thymidylate synthase# or thya

FILE 'MEDLINE'

4825 THYMIDYLATE

113608 SYNTHASE#

3978 THYMIDYLATE SYNTHASE#

(THYMIDYLATE (W) SYNTHASE#)

180 THYA

L1 4077 THYMIDYLATE SYNTHASE# OR THYA

FILE 'SCISEARCH'

5729 THYMIDYLATE

138898 SYNTHASE#

4456 THYMIDYLATE SYNTHASE#

(THYMIDYLATE(W)SYNTHASE#)

111 THYA

L2 4506 THYMIDYLATE SYNTHASE# OR THYA

FILE 'LIFESCI'

1313 "THYMIDYLATE"

33454 SYNTHASE#

963 THYMIDYLATE SYNTHASE#

("THYMIDYLATE" (W) SYNTHASE#)

103 THYA

L3 1016 THYMIDYLATE SYNTHASE# OR THYA

FILE 'BIOTECHDS'

232 THYMIDYLATE

7582 SYNTHASE#

177 THYMIDYLATE SYNTHASE#

(THYMIDYLATE(W)SYNTHASE#)

60 THYA

L4 218 THYMIDYLATE SYNTHASE# OR THYA

FILE 'BIOSIS'

6055 THYMIDYLATE

125880 SYNTHASE#

3757 THYMIDYLATE SYNTHASE#

(THYMIDYLATE(W)SYNTHASE#)

185 THYA

L5 3874 THYMIDYLATE SYNTHASE# OR THYA

FILE 'EMBASE'

4747 "THYMIDYLATE"

114866 SYNTHASE#

4064 THYMIDYLATE SYNTHASE#

("THYMIDYLATE" (W) SYNTHASE#)

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134 THYA
          4135 THYMIDYLATE SYNTHASE# OR THYA
L6
FILE 'HCAPLUS'
          6316 THYMIDYLATE
        122782 SYNTHASE#
          3887 THYMIDYLATE SYNTHASE#
                 (THYMIDYLATE (W) SYNTHASE#)
           280 THYA
L7
          4058 THYMIDYLATE SYNTHASE# OR THYA
FILE 'NTIS'
            20 THYMIDYLATE
           304 SYNTHASE#
             5 THYMIDYLATE SYNTHASE#
                 (THYMIDYLATE(W)SYNTHASE#)
             2 THYA
             7 THYMIDYLATE SYNTHASE# OR THYA
L8
FILE 'ESBIOBASE'
          1803 THYMIDYLATE
         58549 SYNTHASE#
          1533 THYMIDYLATE SYNTHASE#
                 (THYMIDYLATE (W) SYNTHASE#)
            69 THYA
L9
          1560 THYMIDYLATE SYNTHASE# OR THYA
FILE 'BIOTECHNO'
          1423 THYMIDYLATE
         29457 SYNTHASE#
          1155 THYMIDYLATE SYNTHASE#
                 (THYMIDYLATE (W) SYNTHASE#)
            74 THYA
L10
          1195 THYMIDYLATE SYNTHASE# OR THYA
FILE 'WPIDS'
           291 THYMIDYLATE
          7638 SYNTHASE#
           202 THYMIDYLATE SYNTHASE#
                  (THYMIDYLATE (W) SYNTHASE#)
            42 THYA
L11
           230 THYMIDYLATE SYNTHASE# OR THYA
TOTAL FOR ALL FILES
         24876 THYMIDYLATE SYNTHASE# OR THYA
L12
=> s (biologic? or microorganism?)(10a)(control? or containment or abalat?)
FILE 'MEDLINE'
        862799 BIOLOGIC?
         43223 MICROORGANISM?
       2560565 CONTROL?
          8002 CONTAINMENT
             4 ABALAT?
L13
         13561 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'SCISEARCH'
        431305 BIOLOGIC?
         58817 MICROORGANISM?
       2093908 CONTROL?
          8514 CONTAINMENT
            11 ABALAT?
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```
T.14
         25622 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'LIFESCI'
        172952 BIOLOGIC?
         49464 MICROORGANISM?
        509936 CONTROL?
          1252 CONTAINMENT
             2 ABALAT?
L15
         22965 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'BIOTECHDS'
         63301 BIOLOGIC?
         29908 MICROORGANISM?
         73069 CONTROL?
           426 CONTAINMENT
             0 ABALAT?
L16
          6558 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'BIOSIS'
        508841 BIOLOGIC?
       3371749 MICROORGANISM?
       2464549 CONTROL?
          3754 CONTAINMENT
            20 ABALAT?
L17
         55026 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'EMBASE'
        412230 BIOLOGIC?
        136329 MICROORGANISM?
       3826864 CONTROL?
          5681 CONTAINMENT
             6 ABALAT?
L18
          8724 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'HCAPLUS'
       4038046 BIOLOGIC?
        712020 BIOL
       4445630 BIOLOGIC?
                  (BIOLOGIC? OR BIOL)
        182630 MICROORGANISM?
       2784598 CONTROL?
         14634 CONTAINMENT
             4 ABALAT?
L19
         62707 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'NTIS'
         54339 BIOLOGIC?
          9479 MICROORGANISM?
        347093 CONTROL?
         12884 CONTAINMENT
             5 ABALAT?
L20
          2800 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'ESBIOBASE'
```

207927 BIOLOGIC?

```
753246 CONTROL?
          1473 CONTAINMENT
             0 ABALAT?
L21
         17489 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
FILE 'BIOTECHNO'
        93054 BIOLOGIC?
        18193 MICROORGANISM?
        620701 CONTROL?
           536 CONTAINMENT
            1 ABALAT?
          5894 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
L22
               ABALAT?)
FILE 'WPIDS'
        199726 BIOLOGIC?
         1411 BIOL
        200793 BIOLOGIC?
                 (BIOLOGIC? OR BIOL)
        63214 MICROORGANISM?
       3302531 CONTROL?
         14741 CONTAINMENT
             2 ABALAT?
L23
          9054 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
               ABALAT?)
TOTAL FOR ALL FILES
L24 230400 (BIOLOGIC? OR MICROORGANISM?) (10A) (CONTROL? OR CONTAINMENT OR
              ABALAT?)
=> s 112 and 124
FILE 'MEDLINE'
L25
           5 L1 AND L13
FILE 'SCISEARCH'
L26
           8 L2 AND L14
FILE 'LIFESCI'
            3 L3 AND L15
FILE 'BIOTECHDS'
L28
            6 L4 AND L16
FILE 'BIOSIS'
            5 L5 AND L17
L29
FILE 'EMBASE'
     7 L6 AND L18
L30
FILE 'HCAPLUS'
           17 L7 AND L19
L31
FILE 'NTIS'
L32
            0 L8 AND L20
FILE 'ESBIOBASE'
L33
            4 L9 AND L21
FILE 'BIOTECHNO'
```

160352 MICROORGANISM?

L34

2 L10 AND L22

FILE 'WPIDS'

L35 3 L11 AND L23

TOTAL FOR ALL FILES

L36 60 L12 AND L24

=> s 124 and (thymidine or thymine)

FILE 'MEDLINE'

66784 THYMIDINE

12060 THYMINE

L37 31 L13 AND (THYMIDINE OR THYMINE)

FILE 'SCISEARCH'

32015 THYMIDINE

8449 THYMINE

L38 30 L14 AND (THYMIDINE OR THYMINE)

FILE 'LIFESCI'

13496 THYMIDINE

3113 THYMINE

L39 14 L15 AND (THYMIDINE OR THYMINE)

FILE 'BIOTECHDS'

3834 THYMIDINE

1052 THYMINE

L40 24 L16 AND (THYMIDINE OR THYMINE)

FILE 'BIOSIS'

60004 THYMIDINE

10058 THYMINE

L41 36 L17 AND (THYMIDINE OR THYMINE)

FILE 'EMBASE'

58777 THYMIDINE

9415 THYMINE

L42 30 L18 AND (THYMIDINE OR THYMINE)

FILE 'HCAPLUS'

56856 THYMIDINE

20116 THYMINE

L43 128 L19 AND (THYMIDINE OR THYMINE)

FILE 'NTIS'

552 THYMIDINE

236 THYMINE

L44 5 L20 AND (THYMIDINE OR THYMINE)

FILE 'ESBIOBASE'

12605 THYMIDINE

2390 THYMINE

L45 10 L21 AND (THYMIDINE OR THYMINE)

FILE 'BIOTECHNO'

17232 THYMIDINE

3243 THYMINE

L46 14 L22 AND (THYMIDINE OR THYMINE)

FILE 'WPIDS'

4373 THYMIDINE

2226 THYMINE

L47 19 L23 AND (THYMIDINE OR THYMINE)

TOTAL FOR ALL FILES

L48 341 L24 AND (THYMIDINE OR THYMINE)

=> s (136 or 148) not 2003-2009/py

FILE 'MEDLINE'

4037483 2003-2009/PY

L49 22 (L25 OR L37) NOT 2003-2009/PY

FILE 'SCISEARCH'

7579956 2003-2009/PY

(20030000-20099999/PY)

L50 21 (L26 OR L38) NOT 2003-2009/PY

FILE 'LIFESCI'

1052983 2003-2009/PY

L51 8 (L27 OR L39) NOT 2003-2009/PY

FILE 'BIOTECHDS'

148657 2003-2009/PY

L52 18 (L28 OR L40) NOT 2003-2009/PY

FILE 'BIOSIS'

3618282 2003-2009/PY

L53 27 (L29 OR L41) NOT 2003-2009/PY

FILE 'EMBASE'

3485325 2003-2009/PY

L54 20 (L30 OR L42) NOT 2003-2009/PY

FILE 'HCAPLUS'

7998614 2003-2009/PY

L55 86 (L31 OR L43) NOT 2003-2009/PY

FILE 'NTIS'

104791 2003-2009/PY

L56 5 (L32 OR L44) NOT 2003-2009/PY

FILE 'ESBIOBASE'

2059486 2003-2009/PY

L57 7 (L33 OR L45) NOT 2003-2009/PY

FILE 'BIOTECHNO'

122467 2003-2009/PY

L58 13 (L34 OR L46) NOT 2003-2009/PY

FILE 'WPIDS'

6715892 2003-2009/PY

L59 5 (L35 OR L47) NOT 2003-2009/PY

TOTAL FOR ALL FILES

L60 232 (L36 OR L48) NOT 2003-2009/PY

=> dup rem 160

PROCESSING COMPLETED FOR L60

L61 155 DUP REM L60 (77 DUPLICATES REMOVED)

=> d tot

L61 ANSWER 1 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN

TI Identifying nucleic acid ligands photocrosslinking to target from nucleic acids containing photoreactive groups, by modification of systematic

- evolution of ligands by exponential enrichment method, termed photoSELEX; recombinant basic fibroblast growth factor ligand screening for use in diagnosis
- AU GOLD L; SMITH J D; KOCH T; GOLDEN M
- AN 2002-09653 BIOTECHDS
- PI WO 2002006510 24 Jan 2002
- L61 ANSWER 2 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN
- TI Measuring abundance and expression of indicator and effector genes within biotreatment system, by sampling activated sludges and effecting polymerase chain reaction amplification of indicator and effector gene combinations from sample;

reverse transcription-polymerase chain reaction and DNA primer for activated sludge monitoring

- AU CARSON D B; RICE J F
- AN 2003-07131 BIOTECHDS
- PI WO 2002085791 31 Oct 2002
- L61 ANSWER 3 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN
- TI New cyclic nucleotide phosphodiesterase polypeptides such as PDE8A, PDE7A3, TbPDE2A, TbPDE2B, TbPDE2C or TbPDE2E, that are involved in T cell activation, useful for diagnosis and treatment of immune disorders; recombinant enzyme gene production, vector expression in host cell, antibody, sense, antisense molecule, agonist, antagonist and polymerase chain reaction useful in disease gene therapy, drug screening and vaccine
- AU BEAVO J A; SEEBECK T; SODERLING S H; RASCON A; ZORAGHI R; KUNZ S; GONG K; GLAVAS N
- AN 2002-12822 BIOTECHDS
- PI WO 2002022661 21 Mar 2002
- L61 ANSWER 4 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Papovavirus-derived episomal vector and replication control expression system using a mutant large T antigen for human gene therapy and protein production
- SO U.S. Pat. Appl. Publ., 42 pp., Cont.-in-part of U.S. Ser. No. 935,368. CODEN: USXXCO
- IN Cooper, Mark J.
- AN 2002:833420 HCAPLUS
- DN 137:334050

	PATENT NO.					KIND DATE				APP1	LICAT		DATE					
ΡI	US 20020160516					A1	_	2002	1031		US 2	2002-	4328	 9		2	0020	
	US	6339	065			В1		20020	0115		US :	1996-	5942	99		19	9960:	130
	US 5770374				Α		19980	0623		US :	1996–	7286	8 0		19	9961	010	
	WO 9859059				A1 19981230					WO :	1998 <mark>-</mark> 1	JS12	777		19980619			
		W:	AL,	AM,	ΑT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	, BY,	CA,	CH,	CN,	CU,	CZ,	DE,
			DK,	EE,	ES,	FI,	GB,	GE,	GH,	GM,	GW.	, HU,	ID,	IL,	IS,	JP,	ΚE,	KG,
			KP,	KR,	KΖ,	LC,	LK,	LR,	LS,	LT,	LU,	, LV,	MD,	MG,	MK,	MN,	MW,	MX,
			NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG	, SI,	SK,	SL,	ТJ,	TM,	TR,	TT,
			UA,	UG,	US,	UZ,	VN,	YU,	ZW									
		RW:	GH,	GM,	ΚE,	LS,	MW,	SD,	SZ,	UG,	ZW	, AT,	BE,	CH,	CY,	DE,	DK,	ES,
			FI,	FR,	GB,	GR,	ΙE,	ΙΤ,	LU,	MC,	NL	, PT,	SE,	BF,	ВJ,	CF,	CG,	CI,
			CM,	GΑ,	GN,	ML,	MR,	ΝE,	SN,	TD,	ΤG							
	US 20020031803				A1	20020314				US 2	2001-9		20010824					

- L61 ANSWER 5 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Distinct nongenomic signal transduction pathways controlled by $17\beta\text{-estradiol}$ regulate DNA synthesis and cyclin D1 gene transcription in HepG2 cells
- SO Molecular Biology of the Cell (2002), 13(10), 3720-3729 CODEN: MBCEEV; ISSN: 1059-1524

- AU Marino, Maria; Acconcia, Filippo; Bresciani, Francesco; Weisz, Alessandro; Trentalance, Anna
- AN 2002:818828 HCAPLUS
- DN 138:231897
- L61 ANSWER 6 OF 155 MEDLINE on STN DUPLICATE 2
- TI Role of biological markers in the clinical outcome of colon cancer.
- SO British journal of cancer, (2002 Oct 7) Vol. 87, No. 8, pp. 868-75. Journal code: 0370635. ISSN: 0007-0920.
- AU Nanni O; Volpi A; Frassineti G L; De Paola F; Granato A M; Dubini A; Zoli W; Scarpi E; Turci D; Oliverio G; Gambi A; Amadori D
- AN 2002615329 MEDLINE
- L61 ANSWER 7 OF 155 Elsevier Biobase COPYRIGHT 2009 Elsevier Science B.V. on STN
- AN 2002253271 ESBIOBASE
- TI Role of biological markers in the clinical outcome of colon cancer
- AU Nanni, O.; Volpi, A.; Frassineti, G.L.; De Paola, F.; Granato, A.M.; Dubini, A.; Zoli, W.; Scarpi, E.; Turci, D.; Oliverio, G.; Gambi, A.; Amadori, D.
- CS Nanni, O.; Volpi, A.; Frassineti, G.L.; De Paola, F.; Granato, A.M.; Dubini, A.; Zoli, W.; Scarpi, E.; Turci, D.; Oliverio, G.; Gambi, A.; Amadori, D. (Department of Medical Ontology, Pierantoni Hospital, Via Forlanini 34, 47100 Forli (IT))
- SO British Journal of Cancer (7 Oct 2002) Volume 87, Number 8, pp. 868-875, 63 refs.

 CODEN: BJCAAI ISSN: 0007-0920

 DOI: 10.1038/sj.bjc.6600569
- CY United Kingdom
- DT Journal; Article
- LA English
- SL English
- ED Entered STN: 1 Feb 2009
 Last updated on STN: 1 Feb 2009
- L61 ANSWER 8 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN
- TI Exploitation of genetically modified inoculants for industrial ecology applications;
 - vector-mediated gene transfer and expression in host cell for strain improvement and potential bioremediation or biological control agent
- SO ANTONIE VAN LEEUWENHOEK INTERNATIONAL JOURNAL OF GENERAL AND MOLECULAR MICR; (2002) 81, 1-4, 599-606 ISSN: 0003-6072
- AU MORRISSEY JP; WALSH UF; O'DONNELL A; MOENNE-LOCCOZ Y; O'GARA F
- AN 2002-14785 BIOTECHDS
- L61 ANSWER 9 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- ${\tt TI}$ Controlled aggregation of azobenzene based on DNA-mimetics at the air-water interface
- SO International Journal of Nanoscience (2002), 1(5 & 6), 597-601 CODEN: IJNNAJ; ISSN: 0219-581X
- AU Ijiro, Kuniharu; Matsumoto, Jin; Morisue, Mitsuhiko; Shimomura, Masatsugu
- AN 2004:230627 HCAPLUS
- DN 141:239194
- L61 ANSWER 10 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI RecQ helicases and genome stability: lessons from model organisms and human disease
- SO Swiss Medical Weekly (2002), 132(31/32), 433-442 CODEN: SMWWAI; ISSN: 1424-7860
- AU Bjergbaek, Lotte; Cobb, Jennifer A.; Gasser, Susan M.
- AN 2003:46641 HCAPLUS

- DN 138:269186
- L61 ANSWER 11 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- ${\tt TI}$ Genetic and molecular control of folate-homocysteine metabolism in mutant mice
- SO Mammalian Genome (2002), 13(5), 259-267 CODEN: MAMGEC; ISSN: 0938-8990
- AU Ernest, Sheila; Christensen, Benedicte; Gilfix, Brian M.; Mamer, Orval A.; Hosack, Angela; Rodier, Mitchell; Colmenares, Clemencia; McGrath, James; Bale, Allen; Balling, Rudi; Sankoff, David; Rosenblatt, David S.; Nadeau, Joseph H.
- AN 2002:429880 HCAPLUS
- DN 137:246047
- L61 ANSWER 12 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Identification of Novel E2F1-Regulated Genes by Microarray
- SO Archives of Biochemistry and Biophysics (2002), 399(2), 212-224 CODEN: ABBIA4; ISSN: 0003-9861
- AU Ma, Yihong; Croxton, Rhonda; Moorer, Ronnie L., Jr.; Cress, W. Douglas
- AN 2002:176045 HCAPLUS
- DN 137:120414
- L61 ANSWER 13 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Resveratrol, a chemopreventive agent, disrupts the cell cycle control of human SW480 colorectal tumor cells
- SO International Journal of Molecular Medicine (2002), 10(2), 193-199 CODEN: IJMMFG; ISSN: 1107-3756
- AU Delmas, Dominique; Passilly-Degrace, Patricia; Jannin, Brigitte; Cherkaoui Malki, Mustapha; Latruffe, Norbert
- AN 2002:622327 HCAPLUS
- DN 138:147337
- L61 ANSWER 14 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN
- TI Remote electronic control of DNA hybridization through inductive coupling to an attached metal nanocrystal antenna;
 - DNA synthesis and oligonucleotide immobilization on gold nanocrystal support matrix for molecular study
- SO Nature; (2002) 415, 6868, 152-55 CODEN: NATUAS ISSN: 0028-0836
- AU Hamad-Schifferli K; Schwartz J J; Santos A T; Zhang S; *Jacobson J M
- AN 2001-15626 BIOTECHDS
- L61 ANSWER 15 OF 155 SCISEARCH COPYRIGHT (c) 2009 The Thomson Corporation on STN
- TI Homeostatic control of uridine and the role of uridine phosphorylase: a biological and clinical update
- SO BIOCHIMICA ET BIOPHYSICA ACTA-MOLECULAR BASIS OF DISEASE, (18 JUL 2002) Vol. 1587, No. 2-3, Sp. iss. SI, pp. 133-144. ISSN: 0925-4439.
- AU Pizzorno G (Reprint); Cao D L; Leffert J J; Russell R L; Zhang D K; Handschumacher R E
- AN 2002:586160 SCISEARCH
- L61 ANSWER 16 OF 155 MEDLINE on STN
- TI alpha 2-Macroglobulin: a new component in the insulin-like growth factor/insulin-like growth factor binding protein-1 axis.
- SO The Journal of biological chemistry, (2001 Nov 9) Vol. 276, No. 45, pp. 41668-74. Electronic Publication: 2001-08-23. Journal code: 2985121R. ISSN: 0021-9258.
- AU Westwood M; Aplin J D; Collinge I A; Gill A; White A; Gibson J M
- AN 2001664326 MEDLINE

- L61 ANSWER 17 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Structure Control on Photodimerization of Uracil and Thymine Moieties in Nucleolipid Langmuir-Blodgett Films by the Molecular Recognition Effect at the Air/Water Interface
- SO Langmuir (2001), 17(7), 2228-2234 CODEN: LANGD5; ISSN: 0743-7463
- AU Li, Chun; Huang, Jianguo; Liang, Yingqiu
- AN 2001:156165 HCAPLUS
- DN 134:349526
- L61 ANSWER 18 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Serum stimulation and cell density regulate the proliferation of AsPC-1 cells through control of cyclin E and p27KIP1 expression
- SO Anticancer Research (2001), 21(3B), 1885-1891 CODEN: ANTRD4; ISSN: 0250-7005
- AU Horiguchi-Yamada, Junko; Yoshida, Seiya; Kuhara, Akiko; Aoki, Teruaki; Ohno, Tsuneya; Yamada, Hisashi
- AN 2001:623917 HCAPLUS
- DN 136:197787
- L61 ANSWER 19 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI IGFBPs modulate IGF-I- and high glucose-controlled growth of human retinal endothelial cells
- SO Journal of Endocrinology (2001), 171(2), 273-284 CODEN: JOENAK; ISSN: 0022-0795
- AU Giannini, S.; Cresci, B.; Pala, L.; Ciucci, A.; Franchini, A.; Manuelli, C.; Fujita-Yamaguchi, Y.; Cappugi, P.; Zonefrati, R.; Rotella, C. M.
- AN 2001:849253 HCAPLUS
- DN 136:80319
- L61 ANSWER 20 OF 155 Elsevier Biobase COPYRIGHT 2009 Elsevier Science B.V. on STN
- AN 2001093566 ESBIOBASE
- TI Effect of ionizing radiation on thymidine uptake, differentiation, and VEGFR2 receptor expression in endothelial cells: The role of VEGF 165
- AU Gieschen, Holger L.; Spiro, Ira J.; Suit, Herman D.; Ancukiewicz, Marek; Willett, Christopher G; Ott, Mark J.; Rattner, David W.
- Gieschen, Holger L. (Regional Cancer Center, Cape Cod Hospital, Hyannis, MA (US)); Spiro, Ira J.; Suit, Herman D.; Ancukiewicz, Marek; Willett, Christopher G (Department of Radiation Oncology, Massachusetts General Hospital, Boston, MA (US)); Ott, Mark J. (Department of Surgical Oncology, Massachusetts General Hospital, Boston, MA (US)); Rattner, David W. (Department of General Surgery, Massachusetts General Hospital, Boston, MA (US))
- EMAIL: cwillett@partners.org
- SO International Journal of Radiation Oncology Biology Physics (1 May 2001) Volume 50, Number 1, pp. 213-220, 37 refs.

 CODEN: IOBPD3 ISSN: 0360-3016

 DOI: 10.1016/S0360-3016(01)01445-6
- PUI S0360301601014456
- CY United States of America
- DT Journal; Article
- LA English
- SL English
- ED Entered STN: 1 Feb 2009
 Last updated on STN: 1 Feb 2009
- L61 ANSWER 21 OF 155 BIOTECHNO COPYRIGHT 2009 Elsevier Science B.V. on STN
- TI Effect of ionizing radiation on thymidine uptake, differentiation, and VEGFR2 receptor expression in endothelial cells: The role of VEGF.sub.1.sub.6.sub.5

- SO International Journal of Radiation Oncology Biology Physics, (01 MAY 2001), 50/1 (213-220), 37 reference(s) CODEN: IOBPD3 ISSN: 0360-3016
- AU Kermani P.; Leclerc G.; Martel R.; Fareh J.
- AN 2001:32318579 BIOTECHNO
- L61 ANSWER 22 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Activation of transforming growth factor- $\beta 1$ by hepatic stellate cells in vitro and consequences for cell proliferation and survival
- SO Cells of the Hepatic Sinusoid (2001), 8, 191-194 CODEN: CHSIEL
- AU Williams, E. J.; Cochrane, B. C.; Arthur, M. J. P.; Benyon, R. C.
- AN 2001:630431 HCAPLUS
- DN 135:327722
- L61 ANSWER 23 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Loss of cyclin A and G1-cell cycle arrest are a prerequisite of ceramide-induced toxicity in human arterial endothelial cells
- SO Cardiovascular Research (2001), 50(1), 97-107 CODEN: CVREAU; ISSN: 0008-6363
- AU Spyridopoulos, I.; Mayer, P.; Shook, K. S.; Axel, D. I.; Viebahn, R.; Karsch, K. R.
- AN 2001:219989 HCAPLUS
- DN 135:58957
- L61 ANSWER 24 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Flow-Induced DNA Synthesis Requires Signaling to a Translational Control Pathway
- SO Journal of Surgical Research (2001), 97(1), 20-26 CODEN: JSGRA2; ISSN: 0022-4804
- AU Kraiss, Larry W.; Ennis, Tina M.; Alto, Neal M.
- AN 2001:292743 HCAPLUS
- DN 135:135048
- L61 ANSWER 25 OF 155 BIOSIS COPYRIGHT (c) 2009 The Thomson Corporation on STN
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		IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	KΖ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	
		MD,	MG,	MK,	MN,	MW,	MX,	NO,	NΖ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	
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- ANSWER 8 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN L61 AΒ AUTHOR ABSTRACT - The major growth seen in the biotechnology industry in recent decades has largely been driven by the exploitation of genetic engineering techniques. The initial benefits have been predominantly in the biomedical area, with products such as vaccines and hormones that have received broad public approval. In the environmental biotechnology and industrial ecology sectors, biotechnology has the potential to make significant advances through the use ofgenetically modified (GM) microbial inoculants that can reduce agri-chemical usage or remediate polluted environments. Although many GM inoculants have been developed and tested under laboratory conditions, commercial exploitation has lagged behind. Here, we review scientific and regulatory requirements that must be satisfied as part of that exploitation process. Particular attention is paid to new European Union (EU) regulations (Directives) that govern the testing and release of genetically modified organisms and microbial plant protection inoculants in the EU. With regard to the release ofGM inoculants, the impact of the inoculant and the fate of modified genesare important concerns. Long term monitoring of release sites is necessary to address these issues. Data are reported from the monitoring of a site 6years after release of GM Sinorhizobium meliloti strains. It was found that despite the absence of a host plant, the GM strains persisted in the soilfor at least 6 years. Horizontal transfer and microevolution of a GM plasmid between S. meliloti strains was also observed. These data illustrate theimportance of assessing the long-term persistence of GM inoculants. (8 pages)
- L61 ANSWER 26 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- AB The invention relates to compns. and methods for delivering a virus vector to an animal. The compns. include compns. which comprise a matrix having a virus vector bound at the exterior surface thereof in a physiol. reversible manner. The invention also includes methods of making such compns., including particles, devices, bulk materials, and other objects which comprise, consist of, or are coated with such compns. Methods of delivering a virus vector to an animal tissue are also described.
- L61 ANSWER 27 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN
- AB The invention relates to compns. and methods for delivering a virus vector to an animal. The compns. include compns. which comprise a hydrogel matrix (e.g. a collagen matrix which can comprise a poloxamer or an alginate) containing a virus vector therein in a transfectious form. The invention also includes methods of making such hydrogel precursor mixts. and hydrogel matrixes, including particles, devices, bulk materials, and other objects which comprise, consist of, or are coated with such mixts. or matrixes. The invention further relates to compns. comprising a

hydrogel precursor mixture having a virus vector suspended therein, which, when administered to an animal, gel to form a hydrogel matrix containing a virus vector therein in a transfectious form. Methods of delivering a virus vector to an animal tissue are also described.

- ANSWER 39 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN L61 AΒ A new genetically engineered Pseudomonas sp. biological control agent that can control attacks of crop plants by pathogenic fungi e.g. Rhizoctonia sp. and Pythium sp., and aggressively compete with indigenous bacteria and microflora in the plant rhizosphere, is produced from Pseudomonas fluorescens parent strains modified using lemA and gacA regulatory genes and/or genes involved in the synthesis of the fungicide metabolite phenazine-1-carboxylic acid and/or pyrrolnitrin (PN) to enhance production of the fungicide metabolites. The strains can be applied to e.g. cotton (Gossypium hirsutum), wheat (Triticum aestivum) and bean crop plants, seeds or soil. The preferred strains are as follows: gacA and lemA regulatory genes are transformed into transposon mutant of wild-type P. fluorescens; the first base in gacA is changed from thymidine to adenine; genes involved in PN synthesis are linked to a strong constitutive bacterium promoter; the strain is transformed with a plasmid containing lemA and gacA, optionally with a pmATCD cluster; the strain is transformed with a mutant gacA gene, etc. (others specified). (85pp)
- ANSWER 54 OF 155 HCAPLUS COPYRIGHT 2009 ACS on STN This application relates to tetracycline-controlled eukaryotic expression vectors adapted for use in gene therapy or gene immunization having pos. feedback regulation. The vector constructs comprise a single transcription unit comprising a first cistron encoding a desired gene product and a second cistron encoding the tracycline-controlled activator, and an internal ribosome entry site positioned between the cistrons. Depending on the configuration of the tetracycline-controlled activator-responsive promoter, tetracycline can be used to induce or inhibit transcription. By adjusting the position of the tetracycline operator sequence in relation to the TATA box, the resultant promoter can be modified to function in either pos. or neg. regulation by tetracycline. This general invention is exemplified in a hCMV-IE based plasmid vector. Plasmid pUHD10-3 containing the chimeric tetracycline operator sequence (tetO)/hCMV-IE TATA box, and pUHD15-1, expressing the tetracycline-controlled activator (tTA) were used to construct the plasmid vector. The tTA can be a fusion protein for example one that is modified so that it is localized to the eukaryote nucleus.
- ANSWER 82 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN L61 Liquid-culture parameters (pH, temperature and C-source) were manipulated to AB control phenazine-1-carboxylic acid (PCA) production by the take-all biological control agent Pseudomonas fluorescence 2-79 NRRL B-15132. The optimized fermentor medium contained (per 1) 2 g K2HPO4, 2 g KH2PO4, 0.03 g cytosine, 0.01 g adenine, 0.01 g thymine, 4.4 mg ZnSO4.7H2O, 11 mg CaCl2.2H2O, 10 mg MnCl2.4H2O, 2 mg(NH4)6Mo7024.4H2O, 2.4 mg H3BO3, 0.05 g EDTA, 0.05 mg folic acid, 0.05 mg biotin, 0.05 mg cyanocobalamin, 0.1 g MgSO4.7H2O, 0.01 g NaCl, 0.01 g FeSO4.7H2O and C- and N-sources. Concentrated stock solutions of buffer, Mg2+/Na+, Fe2+, trace minerals, purines/pyrimidines and vitamins were prepared to compose 0.8, 0.2, 1, 1.5, 0.6 and 2% of the total medium volume, respectively. Controlled-pH (7 and 8) studies of C- and N-source utilization for growth and PCA production were carried out in a 2 $\scriptstyle 1$ working volume fermentor at 25 deg, 1,000 ml/min air flow and 300 rpm. High, moderate or low PCA productivities were observed at 25-27, 29-32.5 or 34 deg, respectively. PCA accumulation per unit biomass reached 0.31 g/g on glucose, 0.16 g/g on glycerol and xylose and 0.09 g/g on fructose. (23 ref)

ANSWER 92 OF 155 NTIS COPYRIGHT 2009 NTIS on STN There are different possibilities of biological containment to restrict and to prevent the dispersal of bacterial recombinant DNA after the deliberate release: The use of plasmids with a confined host-range neither transferable by conjugation nor mobilisable, has to be mentioned in this contents. Binding of the recombinant DNA to a specific host is more efficient by integrating the genetical material into the bacterial chromosome. To prevent the dispersal of the organism itself mutations are introduced that lower the competitiveness with the natural bacterial flora leading to a long-term elimination. Another way to eliminate the deliberate released bacteria is the inducible expression of lethal genes. There are some serious problems with the formation of resistant bacteria and effective practical use of the inducing molecules. Considering this none of the introduced systems can said to be a practicable biological containment-mechanism. Genetically modified poxviruses are used for the control of rabies. In this case the biological containment is based on the inactivation of the thymidin-kinase-gene (tk), which is important for the virulence of the virus. The inactivation of the tk-gene is due to the insertion of the glycoprotein G-gene of the rabies-virus. This leads to the expression of the glycoprotein G-gene inducing the production of antibodies against the rabies virus. Baculoviruses are used to control insect pest and valid to have highly restricted host ranges. The biological containment is based on the reduction of the ecological fitness by inactivation of the polyhedrin-codinggene. Biological containment of plants (Nicotiana tabacum, Brassica napus) is tried to be achieved by the induction of male sterility. The cells of the tapetum are destroyed by the induction of ribonuclease genes (TA29-RNase T1 or TA29-Barnase). (orig.). (Copyright (c) 1995 by FIZ. Citation number 95:003492.)

L61 AB

- L61 ANSWER 103 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN AΒ Nonantibiotic and autoselection markers for genetically engineered mi croorganism (GEM) vector construction include bialaphos or glyphosate herbicide resistance, resistance to mercuric salts and organomercuri als (useful for GEMs used for cleanup purposes), growth on lactose an d autoselection in thyA, asd or ssb strains. Many vectors capable of replicating in Gram-negative bacteria are based on IncQ, IncPl and I ncW replicons. However, nearly all broad-host-range plasmids carry a ntibiotic-resistance markers, are unstable in the absence of selective pressure and cause physiological stress. The use of transposons in stead of plasmids as vectors may overcome these problems. A series o f transposon Tn5 and transposon Tn10 derived minitransposons vectors containing nonantibiotic selection determinants has been developed. Expression of recombinant genes in the field can be regulated by mani pulating expression to be affected by a signal present in the contami nated location, or by the use of stationary-phase promoters. Systems are being developed for the biological containment of GEMs in the en vironment. (83 ref)
- ANSWER 104 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN

 There is considerable potential for the genetic manipulation both of the biosynthesis and uptake of siderophores and the production of fungicides by disease-suppressive pseudomonad strains. Use of constitutive siderophore-producing strains, the acquisition of additional ferric siderophore receptor genes, and transfer of fungicide biosynthetic genes to nonproducers offer ways to improve the capabilities of inoculant strains. The development of a stable vector system is a prerequisite for environmental release of genetically engineered Pseudomonas spp. The thymidylate-synthase (TS, EC-2.1.1.45) gene (

thyA) of Lactococcus lactis has been used as a positive selectable marker in various microorganisms. The thy system is based on a host strain which is deficient in TS activity and a vector containing a copy of the L. lactis thyA gene. Since TS activity is essential for de novo DNA synthesis, the vector containing a copy of the gene is stably maintained. The thy system has been demonstrated in Rhizobium meliloti but has not yet been successfully adapted to Pseudomonas sp. (25 ref)

L61 ANSWER 126 OF 155 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN AΒ The possibility of selecting double revertants of Thy+Tdr in Bacillus thuringiensis was investigated. Bac. thuringiensis var. galleriae 351 and mutant derivatives thy (thymine) dra (deoxyriboaldolase) and thy drm (phosphodeoxyribomutase) were studied. There was a variation in the phenotype of the rough colony morphology (R) Thy+ thymidine resistant (Tdr)-forms selected by 4 different methods; thymine prototrophy, resistance to thymidine, phage Tg4 and tetracycline. R Strains could be selected during the selection of Thy+ Tdr variants. The drm genes behaved like the dra gene, although a strain with the control marker preserved in the R form was not obtained, so that the observed reversion of the drm gene during the S to R transformation was not conclusively established. The genetic determinants responsible for antibiotic resistance may be involved in the regulation of the activity of this additional genetic material. (18 ref)

=> fil .becpat COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 390.81 391.03 DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL. ENTRY SESSION CA SUBSCRIBER PRICE -2.46-2.46

FILES 'BIOTECHDS, HCAPLUS, WPIDS' ENTERED AT 22:18:17 ON 15 MAR 2009 ALL COPYRIGHTS AND RESTRICTIONS APPLY. SEE HELP USAGETERMS FOR DETAILS.

3 FILES IN THE FILE LIST

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39684 WO/PC

33881 PRY<=2002

(PRY <= 2002)

148621 PY>=2003

(PY > = 2003)

L62 2 (L28 OR L40) AND WO/PC AND PRY<=2002 AND PY>=2003

FILE 'HCAPLUS'

451994 WO/PC

784236 PRY<=2002

7587616 PY>=2003

L63 4 (L31 OR L43) AND WO/PC AND PRY<=2002 AND PY>=2003

FILE 'WPIDS'

829294 WO/PC

1647855 PRY<=2002

5628701 PY>=2003

(PY > = 2003)

L64 2 (L35 OR L47) AND WO/PC AND PRY<=2002 AND PY>=2003

TOTAL FOR ALL FILES

L65 8 (L36 OR L48) AND WO/PC AND PRY<=2002 AND PY>=2003

=> dup rem 165

PROCESSING COMPLETED FOR L65

L66 7 DUP REM L65 (1 DUPLICATE REMOVED)

=> d tot

- L66 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Sequences of human and mouse telomerase reverse transcriptase (TERT) promoter and their uses in driving expression of therapeutic gene and drug screening
- SO U.S., 53 pp., Cont.-in-part of U.S. Ser. No. 974,584. CODEN: USXXAM
- IN Morin, Gregg B.; Lichtsteiner, Serge P.; Vasserot, Alain P.; Adams, Robert
 R.; Andrews, William H.
- AN 2004:669756 HCAPLUS
- DN 141:200156

DN	PATENT NO.							DATE			APPLICATION NO.						ATE		
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	CA	2362	367			С		2004	0803										
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			SK,	SL,	ТJ,	TM,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZW		
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	US 20060281106								4 US 2006-411604						20060425 <				
	US 7378244														20080425 <				
	US 20080220438				A1		2008	0911		US 2	-800	1096	15		2	J080.	425 <	<	

- L66 ANSWER 2 OF 7 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN
- TI Regulating production of a product in a cell, comprises inserting a regulatable catalytically active nucleic acid into a gene that produces the product or regulates the production of the product in the cell; vector-mediated reporter gene transfer and expression in host cell for

vector-mediated reporter gene transfer and expression in host cell for gene therapy

- AU WILSON C; CLOAD S T; KEEFE A D
- AN 2003-14785 BIOTECHDS
- PI WO 2003027310 3 Apr 2003
- L66 ANSWER 3 OF 7 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN
- TI New human anti-MUC18 monoclonal antibodies, useful for treating a disease or condition associated with expression of MUC18 in a patient, e.g. tumors, cancers, and other malignancies;

vector-mediated gene transfer and expression in host cell and mouse hybridoma cell culture for monoclonal antibody production for use in cancer therapy

- AU GUDAS J
- AN 2003-21745 BIOTECHDS
- PI WO 2003057838 17 Jul 2003
- L66 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2009 ACS on STN
- TI Mucoadhesive erodible drug delivery device for controlled administration of pharmaceuticals and other active compounds
- SO PCT Int. Appl., 46 pp. CODEN: PIXXD2
- IN Moro, Daniel G.; Callahan, Howard; Nowotnik, David P.
- AN 2003:154225 HCAPLUS
- DN 138:210299

DN	138:210299 PATENT NO.									APPLICATION NO.						DATE				
PI	WO	2003 2003	0157	48		A2		2003	0227		WO 2	002-	US26	083		2	0020	816	<	
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		NZ 531766										002-								
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	_	2343		0.1		-		20090120			-			-		20020816 <				
		MX 2004001491 ZA 2004002067								7 MX 2004-1491 8 ZA 2004-2067										

- L66 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2009 ACS on STN
- ${\tt TI}$ Method for identifying cellular targets using reporter constructs under the control of a enhancer or silencer
- SO U.S. Pat. Appl. Publ., 15 pp. CODEN: USXXCO
- IN Erives, Albert J.; Starr, D. Barry
- AN 2003:892334 HCAPLUS
- DN 139:359906

	PATENT NO.					KIND DAT			DATE AP			APPLICATION NO.					ATE	
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	WO 2005078069			A1 20050825		WO 2003-US14788						2	0030	509 <				
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     AU 2003304710
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                                 20050901 AU 2003-304710
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    ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2009 ACS on STN
L66
     Sequences of smooth muscle myosin heavy chain promoter/enhancer for
ΤI
     expressing polynucleotides specifically in smooth muscle cells in vivo
     U.S. Pat. Appl. Publ., 75 pp., Cont.-in-part of U.S. Ser. No. 600,319.
SO
     CODEN: USXXCO
ΤN
     Owens, Gary K.; Manabe, Ichiro
     2003:58708 HCAPLUS
AN
DN
     138:132218
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     PATENT NO.
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                                             US 2002-57726
     US 20030017549
                          A1
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                                                                     20020124 <--
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     US 6914136
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                                 20050705
     WO 9936101
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     US 6780610
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                                 20040824 US 2000-600319
                                                                      20000713 <--
L66 ANSWER 7 OF 7 WPIDS COPYRIGHT 2009
                                                 THOMSON REUTERS on STN
     Measuring abundance and expression of indicator and effector genes within
     biotreatment system, by sampling activated sludges and effecting
     polymerase chain reaction amplification of indicator and effector gene
     combinations from sample
PΙ
     WO 2002085791
                     A2 20021031 (200305)* EN 47[8]
         RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ
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             KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT
             RO RU SD SE SG SI SK SL TJ TM TN TR TT TZ UA UG US UZ VN YU ZA ZM
     US 20030092020 A1 20030515 (200335) EN
     EP 1414750 A2 20040506 (200430) EN
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          R: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT LU LV MC MK NL PT
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     AU 2002257205
                     A1 20021105 (200433) EN
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                     A1 20040201 (200473)
     MX 2003009732
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                      A 20040922 (200503)
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                      B2 20050201 (200511)
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     IN 2003CN01665 P4 20051125 (200607)
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     CN 1268767
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     AU 2002257205 B2 20060921 (200712)
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     CARSON D B; RICE J F
ΙN
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NOVELTY - Regulating production of a product in a cell, is new.

DETAILED DESCRIPTION - Regulating production of a product in a cell comprises inserting a regulatable catalytically active nucleic acid (RCANA) into a gene that produces the product or regulates the production of the product in the cell, where the RCANA comprises a catalytic domain which modifies a transcript to alter its coding potential and a regulatory domain that recognizes an effector that alters the function of the catalytic domain, and contacting the regulatory domain with an effector to regulate production of the product. INDEPENDENT CLAIMS are also included for the following: (1) regulating a biological pathway in cell; and (2) screening a population of cells for a cell that produces a bioproduct.

WIDER DISCLOSURE - Also disclosed are the following: (1) isolating a regulatable catalytically active nucleic acid (RCANA) created by randomizing at least one nucleotide in the catalytic domain of a catalytically active nucleic acid to create a nucleic acid pool; (2) modulating expression of a nucleic acid by providing a polynucleotide that is regulated by a peptide; and (3) an RCANA construct with a regulatable oligonucleotide sequence having a regulatory domain.

BIOTECHNOLOGY - Preferred Method: In regulating production of a product in a cell, the production of the product is fully inhibited, or increased compared to a normal control level. The production of the product is partially inhibited according to the concentration of the effector. The concentration of the effector modulates the activity of the catalytic domain of the regulatable catalytically active nucleic acid (RCANA), where the RCANA blocks or activates the expression of the gene. The effector is the product, a feedback inhibitor of the gene, or an intermediate in a metabolic pathway. The product is produced in a metabolic pathway that is being regulated, or is an intermediate in a metabolic pathway. The biological pathway is preferably a metabolic pathway. The effector is endogenous or exogenous to the cell. The effector or the product is a protein, an enzyme, a protein pharmaceutical, a metabolite, a drug, a dye, a vitamin, a food additive, a chemical additive, a pesticide, an insecticide, a feed compound, or a waste product. The drug is an antibiotic, an anticancer drug, an antifungal, a cholesterol-lowering drug, or an immunosuppressant. Preferably, the effector is an endproduct of a biosynthetic process. Regulating a biological pathway in cell comprises: (a) inserting a first RCANA into a first gene that produces a first product or regulates the production of the first product in the biological pathway in a cell, where the first RCANA comprises a catalytic domain which catalyzes cleavage of the RCANA or excision of the RCNA from gene in which it is inserted followed by ligation of the gene at 5' and 3' ends of cleavage site, and a regulatory domain which recognizes an effector that activates a function of the catalytic domain; (b) inserting a second RCANA into a second gene that produces a second product or regulates the production of the second product in the biological pathway in the cell, where the second RCANA comprises a catalytic domain which catalyzes cleavage of the RCANA or excision of the RCNA from gene in which it is inserted followed by ligation of the gene at 5' and 3' ends of cleavage site, and a regulatory domains which recognizes an effector that activates a function of the catalytic domain; and (c) contacting the first regulatory domain with a first effector to regulate production of the first product, and contacting the second regulatory domain with a second effector to regulate production of the second product. The combination of the first and second effectors controls the flux of metabolites through the biological pathway. The biological pathway is a biosynthetic or a metabolic pathway. The biological pathway is fully inhibited or partially inhibited according to the concentration of the first and second effectors. The first product is the second effector. The method further comprises inserting a third RCANA into a third gene that

produces a third product or regulates the production of the third product in the biological pathway in the cell, where the third RCANA comprises a catalytic domain which catalyzes cleavage of the RCANA, or excision of the RCANA from gene in which it is inserted followed by the ligation of the gene at 5' and 3' ends of cleavage site, and a regulatory domain which recognizes an effector that activates a function of the catalytic domain. The first and second RCANAs block or activate expression of the first and second gene. Screening a population of cells for a cell that produces a bioproduct comprises inserting an RCANA into a reporter gene in the population of cells, such that the RCANA is regulated by the bioproduct, where expression of the reporter gene indicates the production of the bioproduct by the cell. The method further comprises isolating the cell that produces the bioproduct. The reporter gene is green fluorescent protein, thymidylate synthase, or beta lactamase.

ACTIVITY - None given. No biological data given. MECHANISM OF ACTION - Gene Therapy.

USE - The methods are useful for regulating a biological pathway in cell, or regulating production of a product in a cell. The regulatable catalytically active nucleic acids (RCANAs) are useful as regulatory elements to control the expression of genes in a metabolic pathway, or as regulated selectable markers to increase a selective pressure favoring or disfavoring production of a targeted bioproduct. The RCANAs are also useful for in vitro or in vivo sensing or detection, and in gene therapy.

EXAMPLE - No relevant example given. (128 pages)

ANSWER 3 OF 7 BIOTECHDS COPYRIGHT 2009 THOMSON REUTERS on STN DERWENT ABSTRACT:

 ${\tt NOVELTY}$ - An isolated monoclonal antibody comprising a heavy chain amino acid or a heavy chain variable domain, where the antibody binds to MUC18, is new.

DETAILED DESCRIPTION - An isolated monoclonal antibody comprising a heavy chain amino acid comprising an amino acid sequence selected from 10 fully defined sequences of 117-123 amino acids, as given in the specification and a heavy chain variable domain encoded by a nucleic acid molecule comprising a sequence selected from 10 fully defined sequences of 352-370 base pairs, as given in the specification, is new.

WIDER DISCLOSURE - Also disclosed as new are: (1) an isolated nucleic acid encoding the antibody; (2) a vector comprising the nucleic acid; (3) a host cell transformed with the nucleic acid molecule; and (4) producing the antibody.

BIOTECHNOLOGY - Preferred Antibody: The monoclonal antibody is a fully human antibody, and further comprises a light chain amino acid comprising a sequence selected from 10 fully defined sequences of 107-112 amino acids, as given in the specification. The antibody is conjugated to a therapeutic agent, e.g. radioisotope, or to a cytotoxic agent, e.g. ricin. The antibody may further comprise a light chain variable domain encoded by a nucleic acid molecule comprising a sequence selected from 10 fully defined sequences of 322-340 base pairs, as given in the specification.

ACTIVITY - Cytostatic. Human patients with tumors were injected with anti-MUC18 antibody, and at periodic times during the treatment, patients were monitored to determine the progression of the tumor, particularly to monitor growth and metastasis. A tumor patient treated with anti-MUC18 antibodies showed lower levels of tumor growth and metastasis compared to the level of tumor growth and metastasis of tumors in patients treated with control antibodies.

 ${\tt MECHANISM}$ OF ACTION - ${\tt MUC18}$ inhibitor. No biological data given.

USE - The monoclonal antibody is useful for treating a disease or condition associated with the expression of MUC18 on the cell surface, e.g. tumors (e.g. melanoma, esophageal, pancreatic or colorectal tumors),

L66 AB carcinomas (e.g. cervical carcinomas and cervical intraepithelial squamous and glandular neoplasia), and cancers (e.g. colorectal, breast or lung cancer) and other malignancies.

ADMINISTRATION - Dosage is 0.1-50 (0.3-20) mg/kg body weight per day. Administration can be through injection or infusion by intravenous, intraperitoneal, intracerebral, subcutaneous, intramuscular, intraocular, intraarterial, intracerebrospinal, intralesional routes, inhalation, or by sustained systemic release.

EXAMPLE - Monoclonal antibodies against MUC18 were developed by sequentially immunizing XenoMouse mice. Initial immunization was with 5 to the power of 6 SK-MEL-28 cells admixed with Complete Freund's Adjuvant. Subsequent boosts were made first with 5 to the power of 6 SK-MEL-28 cells with Incomplete Freund's adjuvant (IFA), followed by 4 injections with 5 microgram of soluble MUC18-human IgG2 Fc fusion protein admixed with IFA, then a final boost of 10 microgram soluble MUC18-human IgG2 Fc fusion protein without adjuvant. Each mouse was immunized either at the base of the tail by intraperitoneal injection or via hind footpad injection with MUC18 recombinant antigen followed by generation of a large number of candidate monoclonal antibodies. Animals were immunized on days 0, 4, 7, 10, 14, 17 and 20; and 4 days later, fusions were performed. After fusion, cells were resuspended in Dulbecco's modified Eagle medium (DMEM), 15 % fetal calf serum (FCS) containing HAT (hypoxanthine, aminopterin and thymidine), and supplemented with L-glutamine, pen/strep, OPI (oxaloacetate, pyruvate, bovine insulin) for culture at 37 degrees centigrade. Cells were plated on 96-well tissue culture plates, and maintained in HAT supplemented media for 2 weeks. Hybridomas were selected and screened for antigen reactivity by Enzyme-Linked Immunosorbent Assay (ELISA). Cloning was performed on selected antigen-positive wells using limited dilution plating. Assay results identified the following anti-MUC18 antibodies: c3.19.1, c6.11.3, c3.10, c3.22, c3.27, c3.45, c3.65, c6.1, c6.9, c6.2 and c6.12. (78 pages)

ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2009 ACS on STN

The present invention relates to a layered pharmaceutical delivery device for the administration of pharmaceuticals or other active compds. to mucosal surfaces. The device may also be used by itself without the incorporation of a therapeutic. The device of the present invention consists of a water-soluble adhesive layer, a non-adhesive, bioerodible backing layer and one or more pharmaceuticals if desired in either or both layers. Upon application, the device adheres to the mucosal surface, providing protection to the treatment site and localized drug delivery. The "Residence Time", the length of time the device remains on the mucosal surface before complete erosion, can be easily regulated by modifications of the backing layer.

L66 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2009 ACS on STN The present invention is directed to nucleic acid constructs and their use AΒ in identifying cellular factors that function in various cellular processes involving gene expression. Such factors include those that participate in signaling pathways to regulate cellular gene expression. These factors may be the targets of known therapeutic agents, novel targets for a test compound, or amenable to altered expression to modulate cellular processes. In a particular embodiment, luciferase reporter construct containing luciferase gene under the control of a PSA regulatory module operably linked to a Simian Virus 40 (SV40) basal promoter, IRES and hygromycin resistance is co-expressed with vectors expressing a prostate cDNA expression library in an androgen dependent prostate cell line for screening pos. or neg. regulatory mols. in the bicalutamide (androgen receptor antagonist). In another particular embodiment, a ${\tt HSV}$ thymidylate kinase gene can be used to replace hygromycin resistance gene or expressed from a second "control" construct under the control of a

basal SV40 promoter, and latter setting is useful for the screening of cDNAs encoding other factors, such as a membrane associated transporter that removes bicalutamide from the cell. In further embodiments, the silencer can be used to replace the PSA regulatory module.

ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2009 ACS on STN

The present invention provides isolated or recombinant polynucleotides which comprise a smooth muscle myosin heavy chain (SM-MHC) promoter/enhancer sequence capable of conferring smooth muscle specific expression in vivo and other regulatory elements of smooth muscle cells (SMC). The invention more particularly relates to methods for the targeted knockout, or over-expression, of genes of interest within smooth muscle cells or within a subtype of smooth muscle cells. The invention further relates to methods of conferring polynucleotide expression in vivo specifically in smooth muscle cells or in subtypes of smooth muscle cells. The invention further provides expression vector comprising SM-MHC promoter/enhancer sequence, genetic engineered host cells comprising an expression vector, and transgenic animals.

L66 ANSWER 7 OF 7 WPIDS COPYRIGHT 2009 THOMSON REUTERS on STN AB WO 2002085791 A2 UPAB: 20050903

NOVELTY - Determining the levels of abundance and expression of an indicator and effector gene combination within a biotreatment system (BS), comprises isolating DNA and mRNA from microorganisms from a stream of a BS, determining levels of indicator gene abundance by quantitative polymerase chain reaction (qPCR) analysis of DNA and levels of effector gene abundance by qRT-PCR analysis of mRNA.

DETAILED DESCRIPTION - Determining the levels of abundance and expression of an indicator and effector gene combination within a biotreatment system (BS), comprises isolating DNA and mRNA from microorganisms from a microorganism-containing stream of a BS, determining levels of indicator gene abundance by quantitative polymerase chain reaction (qPCR) analysis of DNA and levels of effector gene abundance by qRT-PCR analysis of mRNA, where indicator and effector genes are same or different.

INDEPENDENT CLAIMS are also included for the following:

- (1) optimizing a waste treatment system which comprises sampling wastewater from a waste treatment system, collecting solids from the sample, isolating DNA and RNA from the solids, performing qPCR or competitive qPCR on the DNA to determine indicator gene abundance, performing quantitative RT-PCR (qRT-PCR) on the RNA to determine effector gene expression, where the indicator gene abundance correlates with the active microbial content (AMC) of the sample and the effector gene expression correlates with the active bioremedial content (ABC) of the sample, and the system is perturbed and repeating the steps until the AMC and ABC are within an empirically determined optimal operating range; and
- (2) controlling BS, by sampling a microorganism -containing stream of BS, collecting microorganisms from the sample, isolating DNA and RNA from the microorganisms, determining AMC and ABC value for the sample by qPCR analysis of the DNA or RNA, respectively, or determining a specific bioremedial content (SBC) value for the sample by qPCR analysis of the DNA and qRT-PCR analysis of the RNA, and setting a target AMC, ABC or SBC value for the sample, comparing the determined AMC, ABC or SBC value to the target AMC, ABC or SBC value, and adjusting control processes to make the determined AMC, ABC or SBC values closer to the target values when repeating the above steps.

USE - Useful for determining levels of abundance and expression of an indicator and effector gene combination within a biotreatment system, controlling a biotreatment system and for optimizing a waste treatment system. The treatment system is a continuous flow activated sludge system, sequencing batch reactor system, a packed bed reactor system, immobilized bacteria system, fluidized bed reactor system, trickling filter system, or

a rotating biological contactor system.

ADVANTAGE - The monitoring method is more sensitive than conventional methods and is also more specific as live cells that actively contribute to the degradative potential are assayed. The PCR-based methods allow for accurate, quantitative measurement of both the amount of DNA present for a given indicator gene and levels of expression for the effector gene.

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	ENTRY	SESSION
FULL ESTIMATED COST	48.61	439.64
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
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